Complete Summary

GUIDELINE TITLE

Evaluation and management of patients with heart failure and preserved left ventricular ejection fraction: HFSA 2006 comprehensive heart failure practice guideline.

BIBLIOGRAPHIC SOURCE(S)

Heart Failure Society of America. Evaluation and management of patients with heart failure and preserved left ventricular ejection fraction. J Card Fail 2006 Feb; 12(1):e80-5. [53 references] PubMed

GUI DELI NE STATUS

This is the current release of the guideline.

This guideline updates a previous version: Heart Failure Society of America. Heart Failure Society of America (HFSA) practice guidelines. HFSA guidelines for management of patients with heart failure caused by left ventricular systolic dysfunction--pharmacological approaches. J Card Fail 1999 Dec; 5(4):357-82.

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INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

IDENTIFYING INFORMATION AND AVAILABILITY DISCLAIMER

SCOPE

DISEASE/CONDITION(S)

Heart failure with preserved left ventricular ejection fraction

GUIDELINE CATEGORY

Evaluation Management Treatment

CLINICAL SPECIALTY

Cardiology Family Practice Internal Medicine

INTENDED USERS

Physicians

GUIDELINE OBJECTIVE(S)

To provide recommendations for the evaluation and management of patients with heart failure and preserved left ventricular ejection fraction

TARGET POPULATION

Patients with heart failure with preserved left ventricular ejection fraction

INTERVENTIONS AND PRACTICES CONSIDERED

- 1. Differential diagnosis using echocardiography, electrocardiography, and stress imaging
- 2. Aggressive blood pressure monitoring
- 3. Counseling regarding low-sodium diet
- 4. Diuretic treatment with thiazide or loop diuretic
- 5. Angiotensin receptor blockers
- 6. Angiotensin-converting enzyme inhibitors
- 7. Beta-blocker treatment
- 8. Calcium channel blockers
- 9. Restoration and maintenance of sinus rhythm

MAJOR OUTCOMES CONSIDERED

Mortality and morbidity associated with heart failure and preserved left ventricular ejection fraction

METHODOLOGY

METHODS USED TO COLLECT/SELECT EVIDENCE

Searches of Electronic Databases Searches of Unpublished Data

DESCRIPTION OF METHODS USED TO COLLECT/SELECT THE EVIDENCE

Databases searched included Medline and Cochrane. In addition, the guideline developers polled experts in specific areas for data.

NUMBER OF SOURCE DOCUMENTS

Not stated

METHODS USED TO ASSESS THE QUALITY AND STRENGTH OF THE EVI DENCE

Weighting According to a Rating Scheme (Scheme Given)

RATING SCHEME FOR THE STRENGTH OF THE EVIDENCE

Level A: Randomized, Controlled, Clinical Trials May be assigned based on results of a single trial

Level B: Cohort and Case-Control Studies Post hoc, subgroup analysis, and meta-analysis Prospective observational studies or registries

Level C: Expert Opinion Observational studies – epidemiologic findings Safety reporting from large-scale use in practice

METHODS USED TO ANALYZE THE EVIDENCE

Systematic Review

DESCRIPTION OF THE METHODS USED TO ANALYZE THE EVIDENCE

Not stated

METHODS USED TO FORMULATE THE RECOMMENDATIONS

Expert Consensus

DESCRIPTION OF METHODS USED TO FORMULATE THE RECOMMENDATIONS

The Heart Failure Society of America (HFSA) Guideline Committee sought resolution of difficult cases through consensus building. Written documents were essential to this process, because they provided the opportunity for feedback from all members of the group. On occasion, consensus of Committee opinion was sufficient to override positive or negative results of almost any form or prior evidence.

RATING SCHEME FOR THE STRENGTH OF THE RECOMMENDATIONS

"Is recommended": Part of routine care Exceptions to therapy should be minimized.

"Should be considered": Majority of patients should receive the intervention. Some discretion in application to individual patients should be allowed.

"May be considered": Individualization of therapy is indicated

"Is not recommended": Therapeutic intervention should not be used

COST ANALYSIS

A formal cost analysis was not performed and published cost analyses were not reviewed.

METHOD OF GUIDELINE VALIDATION

Internal Peer Review

DESCRIPTION OF METHOD OF GUIDELINE VALIDATION

The process of moving from ideas of recommendations to a final document includes many stages of evaluation and approval. Every section, once written, had a lead reviewer and 2 additional reviewers. After a rewrite, each section was assigned to another review team, which led to a version reviewed by the Committee as a whole and then the Heart Failure Society of America (HFSA) Executive Council, representing 1 more level of expertise and experience. Out of this process emerged the final document.

RECOMMENDATIONS

MAJOR RECOMMENDATIONS

The strength of evidence (A, B, C) and strength of recommendations are defined at the end of the "Major Recommendations" field.

- Careful attention to differential diagnosis is recommended in patients with heart failure (HF) and preserved left ventricular ejection fraction (LVEF) to distinguish among a variety of cardiac disorders, because treatments may differ. These various entities may be distinguished based on echocardiography, electrocardiography, and stress imaging (via exercise or pharmacologic means, using myocardial perfusion or echocardiographic imaging). See algorithm in Figure 11.1 in the original guideline document for a detailed approach to differential diagnosis. (Strength of Evidence = C)
- Evaluation for the possibility of ischemic heart disease and inducible myocardial ischemia is recommended in patients with HF and preserved LVEF (see National Guideline Clearinghouse [NGC] summary of Heart Failure Society of America (HFSA) guideline <u>Evaluation and Therapy for Heart Failure</u> in the Setting of Ischemic Heart <u>Disease</u>). (Strength of Evidence = C)

- Aggressive blood pressure monitoring is recommended in patients with HF and preserved LVEF (see the NGC summary of HFSA guideline <u>Managing</u> <u>Patients with Hypertension and Heart Failure</u>). (Strength of Recommendation = C)
- Counseling on the use of a low-sodium diet (see NGC summary of HFSA guideline Nonpharmacologic Management and Health Care Maintenance in Patients with Chronic Heart Failure) is recommended for all patients with HF, including those with preserved LVEF. (Strength of Evidence = C)
- Diuretic treatment is recommended in all patients with HF and clinical evidence of volume overload, including those with preserved LVEF. Treatment may begin with either a thiazide or loop diuretic. In more severe volume overload or if response to a thiazide is inadequate, treatment with a loop diuretic should be implemented. Excessive diuresis, which may lead to orthostatic changes in blood pressure and worsening renal function, should be avoided. (Strength of Evidence = C)
- Angiotensin receptor blockers (ARBs) or angiotensin-converting enzyme (ACE) inhibitors should be considered in patients with HF and preserved LVEF.
 - ARBs (Strength of Evidence = B)
 - ACE inhibitors (Strength of Evidence = C)
- ACE inhibitors should be considered in all patients with HF and preserved LVEF who have symptomatic atherosclerotic cardiovascular disease or diabetes and one additional risk factor. (Strength of Evidence = C)

In patients who meet these criteria but are intolerant to ACE inhibitors, ARBs should be considered. (Strength of Evidence = C)

- Beta-blocker treatment is recommended in patients with HF and preserved LVEF who have:
 - Prior myocardial infarction (Strength of Evidence = A)
 - Hypertension (see NGC summary of HFSA guideline <u>Managing Patients</u> with Hypertension and Heart Failure) (Strength of Evidence = B)
 - Atrial fibrillation requiring control of ventricular rate (Strength of Evidence = B)
- Calcium channel blockers should be considered in patients with:
 - Atrial fibrillation requiring control of ventricular rate in whom betablockers have proven inadequate for this purpose because of intolerance. In these patients, diltiazem or verapamil should be considered. (Strength of Evidence = C)
 - Symptom-limiting angina. (Strength of Evidence = A)
 - Hypertension. Amlodipine should be considered. (Strength of Evidence
 C)
- Measures to restore and maintain sinus rhythm should be considered in patients who have symptomatic atrial flutter-fibrillation, but this decision should be individualized. (Strength of Evidence = C)

Definitions:

Strength of Evidence

Level A: Randomized, Controlled, Clinical Trials May be assigned based on results of a single trial Level B: Cohort and Case-Control Studies Post hoc, subgroup analysis, and meta-analysis Prospective observational studies or registries

Level C: Expert Opinion Observational studies – epidemiologic findings Safety reporting from large-scale use in practice

Strength of Recommendations

"Is recommended": Part of routine care Exceptions to therapy should be minimized.

"Should be considered": Majority of patients should receive the intervention. Some discretion in application to individual patients should be allowed.

"May be considered": Individualization of therapy is indicated

"Is not recommended": Therapeutic intervention should not be used

CLINICAL ALGORITHM(S)

None provided

EVIDENCE SUPPORTING THE RECOMMENDATIONS

TYPE OF EVIDENCE SUPPORTING THE RECOMMENDATIONS

The type of supporting evidence is identified and graded for each recommendation (see the "Major Recommendations").

The recommendations are supported by randomized controlled clinical trials, cohort and case-control studies, and expert opinion.

BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS

POTENTIAL BENEFITS

Accurate evaluation and appropriate management of heart failure (HF) and preserved left ventricular ejection fraction (LVEF)

POTENTIAL HARMS

Not stated

QUALIFYING STATEMENTS

QUALIFYING STATEMENTS

It must be recognized that the evidence supporting recommendations is based largely on population responses that may not always apply to individuals within the population. Therefore, data may support overall benefit of 1 treatment over another but cannot exclude that some individuals within the population may respond better to the other treatment. Thus guidelines can best serve as evidence-based recommendations for management, not as mandates for management in every patient. Furthermore, it must be recognized that trial data on which recommendations are based have often been carried out with background therapy not comparable to therapy in current use. Therefore, physician decisions regarding the management of individual patients may not always precisely match the recommendations. A knowledgeable physician who integrates the guidelines with pharmacologic and physiologic insight and knowledge of the individual being treated should provide the best patient management.

IMPLEMENTATION OF THE GUIDELINE

DESCRIPTION OF IMPLEMENTATION STRATEGY

An implementation strategy was not provided.

IMPLEMENTATION TOOLS

Pocket Guide/Reference Cards Slide Presentation

For information about <u>availability</u>, see the "Availability of Companion Documents" and "Patient Resources" fields below.

INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

IOM CARE NEED

Living with Illness

IOM DOMAIN

Effectiveness Patient-centeredness

IDENTIFYING INFORMATION AND AVAILABILITY

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ADAPTATION

Not applicable: The guideline was not adapted from another source.

DATE RELEASED

1999 (revised 2006 Feb)

GUIDELINE DEVELOPER(S)

Heart Failure Society of America, Inc - Disease Specific Society

SOURCE(S) OF FUNDING

Heart Failure Society of America, Inc

GUIDELINE COMMITTEE

Comprehensive Heart Failure Practice Guideline Committee

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FINANCIAL DISCLOSURES/CONFLICTS OF INTEREST

Committee members and reviewers from the Executive Council received no direct financial support from the Heart Failure Society of America (HFSA) or any other source for the development of the guideline. Administrative support was provided by the Heart Failure Society of America staff, and the writing of the document was performed on a volunteer basis by the Committee. Financial relationships that might represent conflicts of interest were collected for all members of the Guideline Committee and of the Executive Council, who were asked to disclose potential conflicts and recuse themselves from discussions when necessary. Current relationships are shown in Table 1.5 of the "Development and Implementation" companion document (see the "Availability of Companion Documents" field).

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This is the current release of the guideline.

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GUIDELINE AVAILABILITY

Electronic copies: Available from the <u>Heart Failure Society of America, Inc. Web site</u>.

Print copies: Available from the Heart Failure Society of America, Inc., Court International - Suite 240 S, 2550 University Avenue West, Saint Paul, Minnesota 55114; Phone: (651) 642-1633

AVAILABILITY OF COMPANION DOCUMENTS

The following are available:

- Heart Failure Society of America. Executive summary: HFSA 2006 comprehensive heart failure practice guideline. J Card Fail 2006 Feb; 12(1):10-38.
- Heart Failure Society of America. Development and implementation of a comprehensive heart failure practice guideline. J Card Fail 2006 Feb; 12(1):e3-9.
- Heart Failure Society of America. Conceptualization and working definition of heart failure. J Card Fail 2006 Feb; 12(1):e10-11.

Electronic copies: Available from the <u>Heart Failure Society of America, Inc. Web site</u>.

• PowerPoint slides. HFSA 2006 comprehensive heart failure guideline.

Electronic copies: Available from the <u>Heart Failure Society of America, Inc.</u> <u>Web site.</u>

The following is also available:

 Heart Failure Society of America. Pocket guide. HFSA 2006 comprehensive heart failure practice guideline.

Electronic copies: Not available at this time.

Print copies: Available from the Heart Failure Society of America, Inc., Court International - Suite 240 South, 2550 University Avenue West, Saint Paul, Minnesota 55114; Phone: (651) 642-1633

PATIENT RESOURCES

None available

NGC STATUS

This NGC summary was completed by ECRI on July 31, 2006. The information was verified by the guideline developer on August 10, 2006.

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